Influenza Surveillance Report

www.infectiousdisease.dhh.la.gov Week 44: 10/28/18 - 11/3/18

Influenza activity is slowly increasing in Louisiana. Influenza B remains the predominant positive rapid result being reported from clinical laboratories around the state. Rhino/Enteroviruses continue to circulate at a very high level and represent >50% of all non-influenza viruses being reported by laboratories around the state.

The Influenza Surveillance Summary Report describes the results of the tracking done by the Louisiana Office of Public Health Infectious Disease Epidemiology Section (IDEpi). This report relies on data supplied by sentinel surveillance sites, including hospital emergency departments (ED), laboratories and physicians' offices. Sentinel sites provide weekly data on Influenza Like Illness (ILI) and/or laboratory confirmed cases.

Taken together, ILI surveillance and laboratory surveillance provide a clear picture of the influenza activity occurring in Louisiana each week. If you have any questions about our surveillance system or would like more information, please contact Julie Hand at 504-568-8298 or julie.hand@la.gov.

ILI is defined as an illness characterized by cough and/or cold symptoms and a fever of 100° F or greater in the absence of a known cause. While not every case of ILI is a case of influenza, the CDC has found that trends in ILI from sentinel sites are a good proxy measure of the amount of influenza activity in an area. For this reason, all states and territories participating in the national surveillance program monitor weekly ILI ratios from their sentinel surveillance sites.

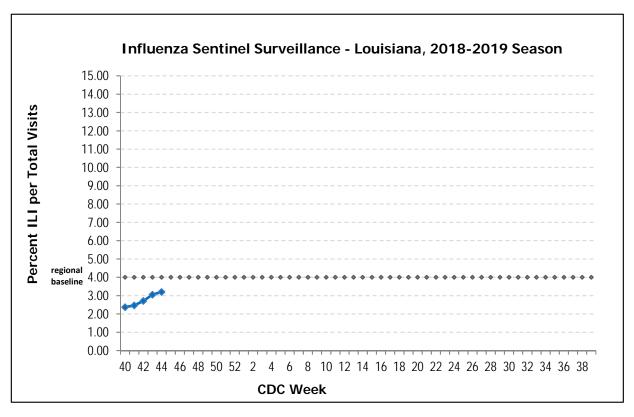


Laboratory testing: Not all sentinel sites have access to laboratory testing. However, many hospitals and physicians' offices do perform some influenza testing. Sites that test for influenza report the number of positive tests each week and the total number of tests performed each week. This information is included on page 3 of this report.

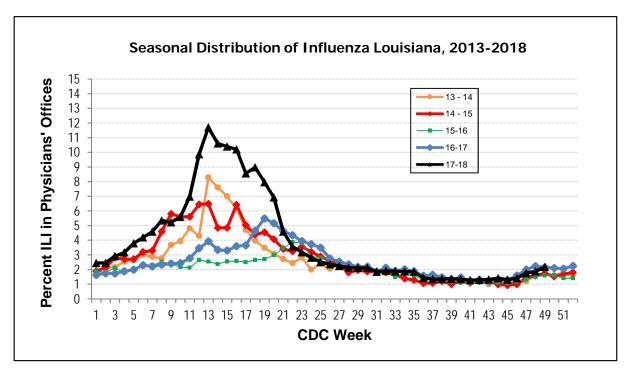
Page 2: ILI Activity

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Page 4: Geographic Distribution
Page 5 & 6: Regional & National Data

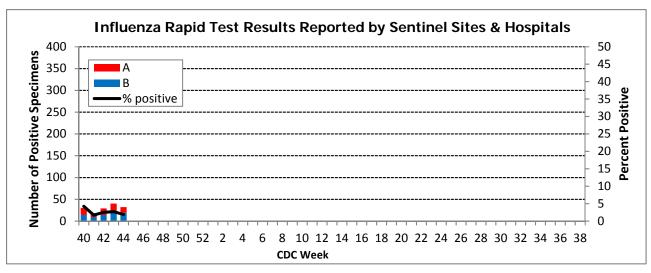
This graph shows the percentage of visits for ILI over the total number of visits for sentinel surveillance sites. This is the best approach to estimate the magnitude of influenza transmission. ILI counts do include some viral infections other than influenza, but experience over the last 50 years has shown that this approach is a reliable method to estimate influenza transmission. It does not show which strain of influenza virus is responsible. The page on lab surveillance does show the proportion of specimens attributable to each virus strain.

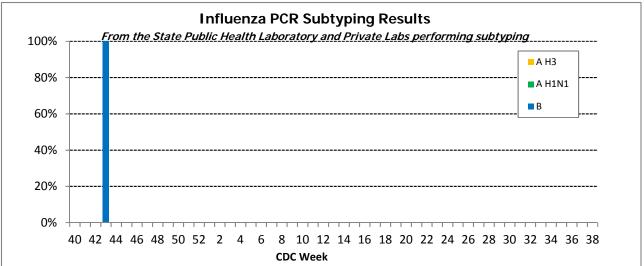


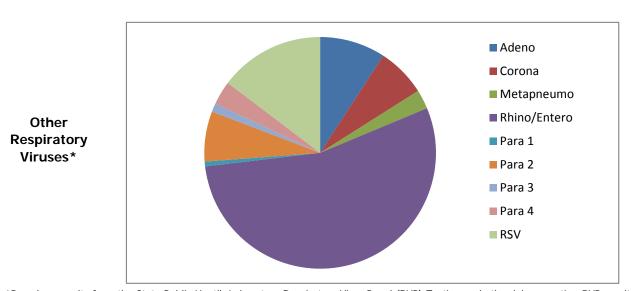
This graph shows the data on ILI surveillance among sentinel physicians' over the past 5 seasons to enable comparisons with previous years and better estimate the amplitude of this season's influenza transmission.



Virologic Surveillance

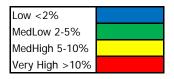






^{*}Based on results from the State Public Heatlh Laboratory Respiratory Virus Panel (RVP) Testing and other labs reporting RVP results over the last 4 weeks.

Geographical Distribution of ILI*



* %IL1 over the last 4 weeks based on sentinel surveillance data

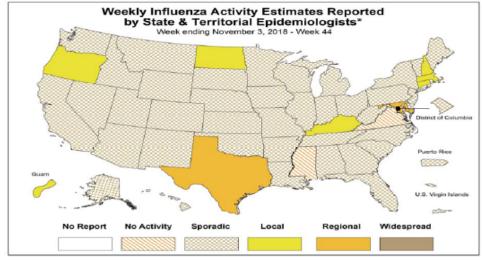


Influenza-Like Illness (ILI) Activity Level Indicator Determined by Data Reported to ILINet 2018-19 Influenza Season Week 44 ending Nov 03, 2018

Geographic Spread of Influenza as Assessed by State and Territorial Epidemiologists







This map indicates geographic spread & does not measure the severity of influenza activity

National Surveillance

Influenza activity in the United States remains low, although small increases in activity were reported.

The proportion of deaths attributed to pneumonia and influenza (P&I) was below the system-specific epidemic threshold.

No influenza-associated pediatric deaths were reported to CDCf for week 44.

The proportion of outpatient visits for influenza-like illness (ILI) was 1.8%, which is below the national baseline of 2.2%.

Clinical Laboratory Data

	Week 44	Data Cumulative since September 30, 2018 (week 40)			
No. of specimens tested	17,264	88,537			
No. of positive specimens (%)	155 (0.9%)	1,538 (1.7%)			
Positive specimens by type					
Influenza A	115 (74.2%)	1,165 (75.7%)			
Influenza B	40 (25.8%)	373 (24.3%)			

Public Health Laboratory Data

	Week 44	Data Cumulative since September 30, 2018 (week 40)
No. of specimens tested	356	3,489
No. of positive specimens*	46	348
Positive specimens by type/subtype		
Influenza A	42 (91.3%)	286 (82.2%)
(H1N1)pdm09	32 (88.9%)	195 (76.8%)
H3N2	4 (11.1%)	59 (23.2%)
Subtyping not performed	6	32
Influenza B	4 (8.7%)	62 (17.8%)
Yamagata lineage	2 (100.0%)	35 (76.1%)
Victoria lineage	0 (0%)	11 (23.9%)
Lineage not performed	2	16

HHS Surveillance Region Data:

Regio	Region 6 (AR, LA, NM, OK, TX)														
CDC	Public Health Labs	Public Health Specimens Tested	AUNK	AH1N1 pdm09	AH3N2	AH3N2v	В	BVic	BYam	Clinical Labs	Clinical Specimens Tested	Clinical Flu Positive	% Positive	A	В
201841	7	73	0	1	0	0	0	1	0	24	2042	32	1.57	16	16
201842	8	69	0	1	0	0	0	0	0	24	2250	37	1.64	19	18
201843	7	53	0	7	2	0	0	0	0	23	2355	28	1.19	16	12
201844	5	20	0	4	1	0	0	0	0	20	3899	24	0.62	17	7

U.S. Out	U.S. Outpatient Influenza-like Illness Surveillance Network (ILINet) 2017-2018 Influenza Season											
HHS	HHS Region 6 (AR, LA, NM, OK, and TX) (Baseline: 4.0%) Data as of Friday, Novembel 9, 2018											
						ILI 65		Total	%	%		
CDC	# Sites	ILI 0-4	ILI 5-24	ILI 25-49	ILI 50-64	and	Total	Patient	Unweighted	Weighted		
Week	Reporting	years	years	years	years	older	ILI	Visits	ILI	ILI		
201841	288	716	676	552	195	103	2242	116453	1.9	2.0		
201842	290	741	787	536	184	138	2386	109749	2.2	2.1		
201843	284	820	932	628	202	133	2715	112931	2.4	2.4		
201844	274	996	945	643	214	152	2950	111645	2.6	2.7		

Antiviral Resistance:

Antiviral Resistance: During May 20-November 3, 2018, 171 specimens (70 influenza A(H1N1)pdm09, 57 influenza A(H3N2), and 44 influenza B viruses) collected in the United States were tested for susceptibility to the neuraminidase inhibitors (oseltamivir, zanamivir, and peramivir). All tested viruses were sensitive to all three recommended antiviral medications.

While all of the recently circulating influenza viruses are susceptible to the neuraminidase inhibitor antiviral medications oseltamivir, zanamivir, and peramivir; rare sporadic instances of oseltamivir-resistant and peramivir-resistant influenza A(H1N1)pdm09 viruses and oseltamivir-resistant influenza A(H3N2) viruses have been detected worldwide. Antiviral treatment as early as possible is recommended for patients with confirmed or suspected influenza who have severe, complicated, or progressive illness; who require hospitalization; or who are at https://www.cdc.gov/flu/antivirals/index.htm.

Antigenic & Genetic Charactization:

Influenza A Viruses

- A (H1N1)pdm09: Phylogenetic analysis of the HA genes from 109 A(H1N1)pdm09 viruses showed that all belonged to clade 6B.1. Seventy A(H1N1)pdm09 viruses were antigenically characterized, and all 70 (100%) were antigenically similar (analyzed using HI with ferret antisera) to A/Michigan/45/2015 (6B.1), a cell-propagated A/Michigan/45/2015-like reference virus representing the A(H1N1)pdm09 component for the 2018-19 Northern Hemisphere influenza vaccines.
- A (H3N2): Phylogenetic analysis of the HA genes from 78 A(H3N2) viruses revealed extensive genetic diversity with multiple clades/subclades co-circulating. The HA genes of circulating viruses belonged to clade 3C.2a (n=28), subclade 3C.2a1 (n=47) or clade 3C.3a (n=3). Fifty-four influenza A(H3N2) viruses were antigenically characterized by FRA with ferret antisera, and 49 (90.7%) A(H3N2) viruses tested were well-inhibited (reacting at titers that were within fourfold of the homologous virus titer) by ferret antisera raised against A/Singapore/INFIMH-16-0019/2016 (3C.2a1), a cell-propagated A/Singapore/INFIMH-16-0019/2016 -like reference virus representing the A(H3N2) component of 2018-19 Northern Hemisphere influenza vaccines.

Influenza B Viruses

- B/Victoria: Phylogenetic analysis of 11 B/Victoria-lineage viruses indicate that all HA genes belonged to genetic clade V1A, however genetic subclades which are antigenically distinct have emerged. The majority of recent B/Victoria-lineage viruses belonged to a subclade of viruses with a 6-nucleotide deletion (encoding amino acids 162 and 163) in the HA (V1A.1, previously abbreviated as V1A-2Del). In addition, a small number of B/Victoria-lineage viruses have a three amino acid deletion (162-164) in the HA protein (abbreviated as V1A-3Del). Nine (90%) B/Victoria lineage viruses were well-inhibited by ferret antisera raised against cell-propagated B/Colorado/06/2017-like (V1A.1) reference virus, representing the B/Victoria lineage component of 2018-19 Northern Hemisphere influenza vaccines. One (10%) B/Victoria lineage virus reacted poorly (at titers that were 8-fold or greater reduced compared with the homologous virus titer) with ferret antisera raised against cell-propagated B/Colorado/06/2017-like reference virus, and belonged to clade V1A.
- B/Yamagata: Phylogenetic analysis of 44 influenza B/Yamagata-lineage viruses indicate
 that the HA genes belonged to clade Y3. A total of 39 influenza B/Yamagata-lineage viruses
 were antigenically characterized, and all were antigenically similar to cell-propagated
 B/Phuket/3073/2013 (Y3), the reference vaccine virus representing the influenza
 B/Yamagata-lineage component of the 2018-19 Northern Hemisphere quadrivalent
 vaccines.